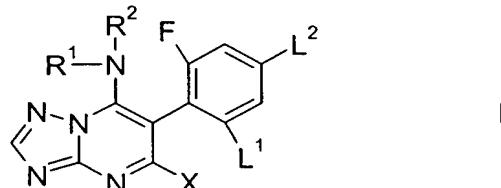


6-(2-Fluorophenyl)-triazolopyrimidines, method for producing them, their use for controlling parasitic fungi and agents containing the same

Abstract

5

6-(2-Fluorophenyl)-triazolopyrimidines of the formula I



in which the substituents are as defined below:

10     R<sup>1</sup> is C<sub>4</sub>-C<sub>8</sub>-alkyl, C<sub>4</sub>-C<sub>8</sub>-haloalkyl, substituted C<sub>3</sub>-C<sub>8</sub>-Cycloalkyl, C<sub>3</sub>-C<sub>8</sub>-halocycloalkyl, C<sub>5</sub>-C<sub>8</sub>-alkenyl, C<sub>2</sub>-C<sub>8</sub>-haloalkenyl, C<sub>3</sub>-C<sub>6</sub>-cycloalkenyl, C<sub>3</sub>-C<sub>6</sub>-halocycloalkenyl, C<sub>2</sub>-C<sub>8</sub>-alkynyl, C<sub>2</sub>-C<sub>8</sub>-haloalkynyl or phenyl, naphthyl, or a five- or six-membered saturated, partially unsaturated or aromatic heterocycle which contains one to four heteroatoms from the group consisting of O, N and S,

15

R<sup>2</sup> is hydrogen, C<sub>1</sub>-C<sub>3</sub>-alkyl or one of the groups mentioned under R<sup>1</sup>,

20     R<sup>1</sup> and R<sup>2</sup> together with the nitrogen atom to which they are attached may also form a five- or six-membered heterocycl or heteroaryl which is attached via N and contain one to three further heteroatoms from the group consisting of O, N and S as ring member,

except for piperidin-1-yl optionally substituted by methyl groups;

25     R<sup>1</sup> and/or R<sup>2</sup> may be substituted according to the description;

L<sup>1</sup> is chlorine or fluorine;

L<sup>2</sup> is hydrogen,

30     is, if L<sup>1</sup> is fluorine, also fluorine;

X is alkyl;

35     processes for preparing these compounds, compositions comprising them and their use for controlling phytopathogenic harmful fungi.